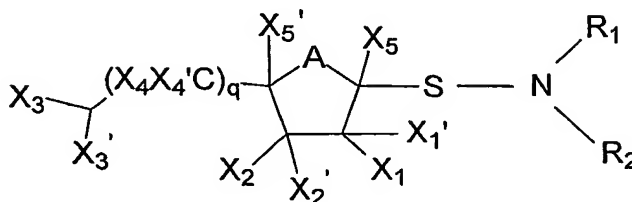


Claims:

1. A compound of general formula (I):



5

wherein A is selected from the group consisting of O, S, SO, SO₂, Se, Te, NR₈, CR₉R'₉, N → O and C(O);

and, when A is O and q is 1, one of R₁ and R₂ is selected from the group consisting of hydrogen, optionally substituted C₁₋₃ or >C₃₀ alkyl, alkyl when interrupted by one or more heteroatoms or functional groups selected from the group consisting of O, S, -N=, NR₇ and -(Y)_mC=(Z)(T)_n-, optionally substituted C₂₋₃ or >C₃₀ alkenyl, alkenyl when interrupted by one or more heteroatoms or functional groups selected from the group consisting of O, S, -N=, NR₇ and -(Y)_mC=(Z)(T)_n-, optionally substituted aralkyl which may be interrupted within the alkyl moiety by one or more heteroatoms or functional groups selected from the group consisting of O, S, -N=, NR₇ and -(Y)_mC=(Z)(T)_n-, optionally substituted acyl, optionally substituted aryl, optionally substituted heterocyclic and a carbohydrate moiety, while the other of R₁ and R₂ is selected from the group consisting of hydrogen, optionally substituted alkyl which may be interrupted by one or more heteroatoms or functional groups selected from the group consisting of O, S, -N=, NR₇ and -(Y)_mC=(Z)(T)_n-, optionally substituted alkenyl which may be interrupted by one or more heteroatoms or functional groups selected from the group consisting of O, S, -N=, NR₇ and -(Y)_mC=(Z)(T)_n-, optionally substituted aralkyl which may be interrupted within the alkyl moiety by one or more heteroatoms or functional groups selected from the group consisting of O, S, -N=, NR₇ and -(Y)_mC=(Z)(T)_n-, optionally substituted

aryl, optionally substituted heterocyclic, optionally substituted acyl and a carbohydrate moiety;

but, when A is S, SO, SO₂, Se, Te, NR₈, CR₉R₉', N → O or C(O) and q is 1 or A is O, S, SO, SO₂, Se, Te, NR₈, CR₉R₉', N → O or C(O) and q is O, then R₁ and R₂ are independently selected from the group consisting of hydrogen, optionally substituted alkyl which may be interrupted by one or more heteroatoms or functional groups selected from the group consisting of O, S, -N=, NR₇ and -(Y)_mC=(Z)(T)_n-, optionally substituted alkenyl which may be interrupted by one or more heteroatoms or functional groups selected from the group consisting of O, S, -N=, NR₇ and -(Y)_mC=(Z)(T)_n-, optionally substituted aralkyl which may be interrupted within the alkyl moiety by one or more heteroatoms or functional groups selected from the group consisting of O, S, -N=, NR₇ and -(Y)_mC=(Z)(T)_n-, optionally substituted aryl, optionally substituted acyl and a carbohydrate moiety, or R₁ and R₂ together with the nitrogen atom from which they depend form a saturated or unsaturated, optionally substituted heterocyclic group which may include additional heteroatoms selected from the group consisting of O, N and S;

X₁ is selected from the group consisting of OR₃, SR₃, NR₃R'₃, hydrogen, halogen, -(Y)_mC=(Z)(T)_nR₃, -N(C=(Z)(T)_nR₃)₂, N₃, CN, OCN, SCN, OSO₃R₃, OSO₂R₃, OPO₃R₃R'₃, OPO₂R₃R'₃, S(O)R₃, S(O)₂R₃, S(O)₂OR₃, PO₃R₃R'₃, NR₃NR'₃R'₃, SNR₃R'₃, NR₃SR'₃, SSR₃ and R₃, or is an oxo group, =S, =NOR₃ or =CR₃R'₃ and X₁' is absent;

X₂ is selected from the group consisting of OR₄, SR₄, NR₄R'₄, hydrogen, halogen, -(Y)_mC=(Z)(T)_nR₄, -N(C=(Z)(T)_nR₄)₂, N₃, CN, OCN, SCN, OSO₃R₄, OSO₂R₄, OPO₃R₄R'₄, OPO₂R₄R'₄, S(O)R₄, S(O)₂R₄, S(O)₂OR₄, PO₃R₄R'₄, NR₄NR'₄R'₄, SNR₄R'₄, NR₄SR'₄, SSR₄ and R₄, or is an oxo group, =S, =NOR₄ or =CR₄R'₄ and X₂' is absent;

X₃ and X₃' are independently selected from the group consisting of OR₅, SR₅, NR₅R'₅, hydrogen, halogen, -

$(Y)_mC=(Z)(T)_nR_5$, $-N(C=(Z)(T)_nR_5)_2$, N_3 , CN , OCN , SCN , OSO_3R_5 , OSO_2R_5 , $OPO_3R_5R'_5$, $OPO_2R_5R'_5$, $S(O)R_5$, $S(O)_2R_5$, $S(O)_2OR_5$, $PO_3R_5R'_5$, $NR_5NR'_5R''_5$, $SNR_5R'_5$, $NR_5SR'_5$, SSR_5 and R_5 , or X_3 is $=O$, $=S$, $=NOR_5$ or $=CR_5R'_5$ and X_3' is absent;

5 X_4 is selected from the group consisting of OR_6 , SR_6 , $NR_6R'_6$, hydrogen, halogen, $-(Y)_mC=(Z)(T)_nR_6$, $-N(C=(Z)(T)_nR_6)_2$, N_3 , CN , OCN , SCN , OSO_3R_6 , OSO_2R_6 , $OPO_3R_6R'_6$, $OPO_2R_6R'_6$, $S(O)R_6$, $S(O)_2R_6$, $S(O)_2OR_6$, $PO_3R_6R'_6$, $NR_6NR'_6R''_6$, $SNR_6R'_6$, $NR_6SR'_6$, SSR_6 and R_6 , or is an oxo group, $=S$, $=NOR_6$ or $=CR_6R'_6$ and X_4' is absent;

10 X_5 is selected from the group consisting of hydrogen, CN , $-C=(Z)(T)_nR_{11}$, $S(O)R_{11}$, $S(O)_2R_{11}$, $S(O)_2OR_{11}$, $PO_3R_{11}R'_{11}$, optionally substituted alkyl, optionally substituted alkaryl, optionally substituted aryl, 15 optionally substituted aralkyl, and optionally substituted acyl;

X_1' , X_2' , X_4' and X_5' are the same or different and are selected from the group consisting of hydrogen, CN , optionally substituted alkyl, optionally substituted 20 alkaryl, optionally substituted aryl, optionally substituted aralkyl, and optionally substituted acyl;

or one of X_1 and X_2 , X_2 and X_5' , X_5' and A when A contains a carbon or nitrogen atom, X_5 and A when A contains a carbon or nitrogen atom, and X_5 and X_1 together 25 constitute a double bond, or X_5' and X_4 or X_3 and X_4 together constitute a double bond, or R_1 and X_1 , R_2 and X_1 , R_1 and X_2 , R_2 and X_2 , R_1 and X_5 , R_2 and X_5 , R_1 and X_5' , R_2 and X_5' , X_1 and X_2 , X_2 and X_3 , X_2 and X_4 , X_3 and X_4 , X_1 and X_1' , X_2 and X_2' , X_3 and X_3' or X_4 and X_4' together form part of a 30 ring structure which optionally includes at least one heteroatom selected from O , S and N and is optionally substituted;

m , n and q are independently 0 or 1 and Y , Z and T are independently selected from the group consisting of 35 O , S , and NR_{10} ;

R_3 , R'_3 , R''_3 , R_4 , R'_4 , R''_4 , R_5 , R'_5 , R''_5 , R_6 , R'_6 , R''_6 , R_7 , R_8 , R_9 , R'_9 , R_{10} , R_{11} and R'_{11} are the same or

different and are selected from the group consisting of hydrogen, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted aryl, optionally substituted aralkyl, optionally substituted acyl and a carbohydrate moiety;

with the proviso that at least two of X_1 , X_2 , X_3 and X_4 are other than hydrogen or a group linked to the ring through a carbon-carbon bond and the further proviso that the compound of general formula (I) is not 1-(9H-purinyl)-S-(3-deoxy-pentafuranosyl)sulfenamide 5 5-formamido-2',3',5'-tri-O-formyl-1-(β -D-ribofuranosylthio)imidazole-4-carboxamide, N-phenyl-S-(2,3:5,6-di-O-isopropylidenyl- β -D-mannofuranosyl)sulfenamide or 10 N,N-diethyl-S-(2,3,5,6-tetra-O-benzoyl- β -D-galactofuranosyl)sulfenamide;

or a pharmaceutically acceptable salt thereof.

2. A compound as claimed in claim 1 wherein q is 0 or q is 1 and A is selected from S, SO, SO₂, Se, Te, NR₈, 20 CR₉R'₉, N \rightarrow O or C(O) and one or both of R₁ and R₂ is alkyl.

3. A compound as claimed in claim 2 wherein one or both of R₁ and R₂ is C₄₋₃₀ alkyl.

4. A compound as claimed in claim 3 wherein one or 25 both of R₁ and R₂ is C₆₋₁₂ alkyl.

5. A compound as claimed in claim 4 wherein one or both of R₁ and R₂ is C₈₋₁₀ alkyl.

6. A compound as claimed in claim 1 wherein one or both of R₁ and R₂ is aralkyl.

7. A compound as claimed in claim 6 wherein one or 30 both R₁ and R₂ is (CH₂)_rPh where Ph is phenyl and r is an integer in the range 1 to 12 inclusive.

8. A compound as claimed in claim 1 wherein one or both of R₁ and R₂ is alkyl interrupted by one or more 35 heteroatoms or functional groups selected from the group consisting of O, S, -N=, NR₇, and -(Y)_mC=(Z)(T)_n.

9. A compound as claimed in claim 8 wherein one or

both of R_1 and R_2 is alkyl interrupted by one or more oxygen atoms.

10. A compound as claimed in claim 9 wherein one or both of R_1 and R_2 is $\text{CH}_3(\text{CH}_2)_x\text{O}(\text{CH}_2)_y\text{O}(\text{CH}_2)_z$ wherein x is an integer in the range 0 to 12 inclusive and y and z are independently integers in the range 1 to 12 inclusive.

11. A compound as claimed in claim 1 wherein q is 0 or q is 1 and A is selected from S , SO , SO_2 , Se , Te , NR_8 , $\text{CR}_9\text{R}'_9$, $\text{N} \rightarrow \text{O}$ or $\text{C}(\text{O})$ and one or both of R_1 and R_2 is alkenyl.

12. A compound as claimed in claim 1 wherein R_1 and R_2 together with the nitrogen atom from which they depend form an optionally substituted saturated or unsaturated heterocyclic group.

13. A compound as claimed in claim 12 wherein R_1 and R_2 together with the nitrogen atom from which they depend form a cyclic imide or a lactam.

14. A compound as claimed in any one of claims 1 to 13 wherein X_1 is OR_3 .

15. A compound as claimed in claim 14 wherein R_3 is hydrogen or optionally substituted acyl.

16. A compound as claimed in any one of claims 1 to 15 wherein X_2 is OR_4 .

17. A compound as claimed in claim 16 wherein R_4 is hydrogen or optionally substituted acyl.

18. A compound as claimed in any one of claims 1 to 17 wherein X_3 is OR_5 .

19. A compound as claimed in claim 18 wherein R_5 is hydrogen or optionally substituted acyl.

20. A compound as claimed in any one of claims 1 to 19 wherein X_4 , when present, is OR_6 .

21. A compound as claimed in claim 20 wherein R_6 is hydrogen or optionally substituted acyl.

22. A compound selected from the group consisting of:

N-benzyl-*S*-(2,3,5,6-tetra-*O*-benzoyl- β -D-galactofuranosyl)sulfenamide

N,N-dibenzyl-*S*-(2,3,5,6-tetra-*O*-acetyl- β -D-galactofuranosyl) sulfenamide

N,N-dicyclohexyl-*S*-(2,3,5,6-tetra-*O*-acetyl- β -D-galactofuranosyl) sulfenamide

5 *N,N*-di(2-methoxyethoxyethyl)-*S*-(2,3,5,6-tetra-*O*-acetyl- β -D-galactofuranosyl) sulfenamide

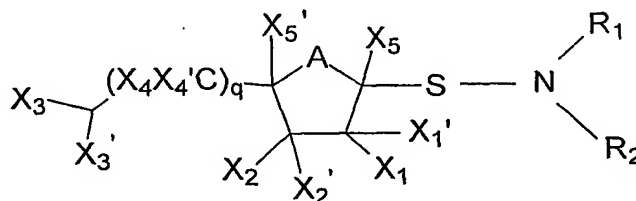
1-(2,2,6,6-tetramethylpiperidiny1)-*S*-(2,3,5,6-tetra-*O*-acetyl- β -D-galactofuranosyl) sulfenamide

10 *N,N*-dioctyl-*S*-(2,3-di-*O*-acetyl-5-*O*-[*tert*-butyldiphenylsilyl]- α -D-arabinofuranosyl) sulfenamide

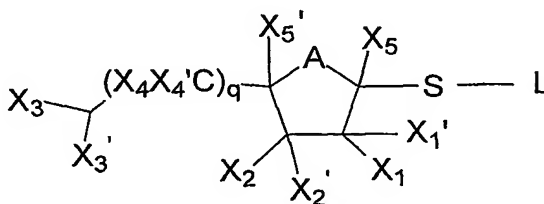
N,N-Dibenzyl-*S*-(β -D-galactofuranosyl) sulfenamide

N,N-Di(2-methoxyethoxyethyl)-*S*-(β -D-galactofuranosyl) sulfenamide

23. A method of preparation of a compound of general
15 formula (I):

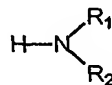


comprising reacting a compound of general formula
20 (II):



wherein L is a leaving group, preferably acetyl
25 and X_1 , X_1' , X_2 , X_2' , X_3 , X_3' , X_4 , X_4' , X_5 and X_5' , are as defined;

with a compound of general formula (III):



wherein R_1 and R_2 are as defined above;
in the presence of a bis-activated alkyl halide.

24. A method for the treatment of a microbial
5 infection, comprising administering to a patient in need
of such treatment a therapeutically effective amount of a
compound of general formula (I) as claimed in any one of
claims 1 to 22.

25. The use of a compound of general formula (I) as
10 claimed in any one of claims 1 to 22 in the manufacture of
a medicament for use in the treatment of a microbial
infection.

26. A pharmaceutical composition comprising a
compound of general formula (I) as claimed in any one of
15 claims 1 to 22 and a pharmaceutically acceptable carrier.

27. A method of killing a microorganism, comprising
exposing said microorganism to a compound of general
formula (I) as claimed in any one of claims 1 to 22.